

INOMax® DS_{IR} Troubleshooting

Warning: Use caution when troubleshooting the INOMax DS_{IR} while in use for a patient. When possible replace the unit in question, and perform troubleshooting procedure once the unit is removed from the patient.

If the system fails to operate properly:


1. Check the patient condition and take appropriate action.
2. Verify that the system is set up as detailed in Section 2/ Setup of the Operation and Maintenance Manual.
3. Use the INOblender® (see INOblender Operation Manual) or backup mode if necessary (see page 29 of manual).
4. Find a symptom or alarm condition in the troubleshooting table which best describes the problem and follow the recommended actions to resolve the problem.



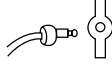


If the problem can't be corrected:



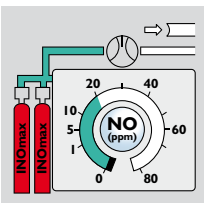
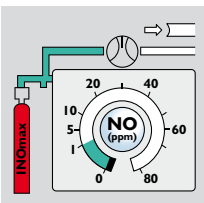
Contact Customer Service on 1300 198 565.

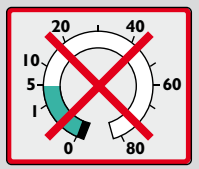
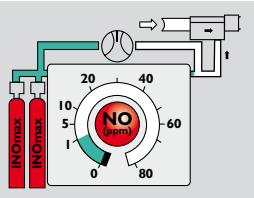
Troubleshooting Guide

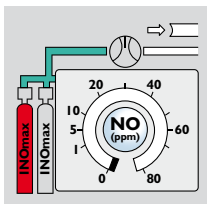
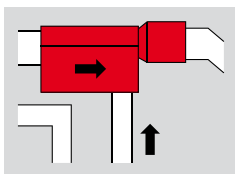
High Priority Alarms		
Symptom/Alarm	Possible Cause	Recommended Action
I. High NO Alarm 	A. Note: A newly installed NO sensor will give high readings until fully conditioned (about 5 hours) and calibrated.	a. After installation of the NO sensor perform a low and high calibration. b. Wait 5 hours and repeat both the low and high calibration.
	B. The High NO alarm level may be inappropriately set.	a. Make sure the High NO alarm is set greater than the Set NO value.
	C. The NO calibration may have drifted.	a. Perform a low and high range calibration of the NO sensor. b. Check calibration sample tee for leaks.
	D. Circuit setup incorrect.	a. Check circuit setup for correct use of adapters and/or check valves.



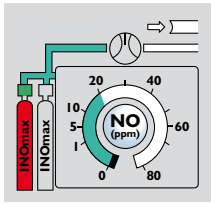
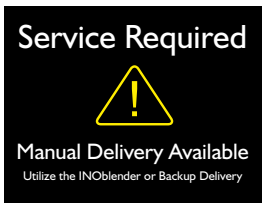
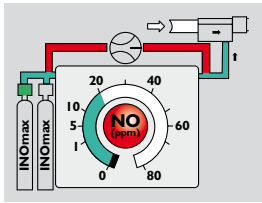
High Priority Alarms		
Symptom/Alarm	Possible Cause	Recommended Action
2. Low NO Alarm  Low NO	A. The Patient Gas Sample line may be disconnected.	a. Reconnect the Patient Gas Sample line.
	B. The Low NO alarm setting may be inappropriately set.	a. Make sure the Low NO alarm is set less than the Set NO value.
	C. The NO calibration may have drifted.	a. Perform a low and high range calibration of the NO sensor.
	D. The NO sensor may not be properly seated.	a. Make sure the sensors are correctly seated with the O-rings and the sensor cover is fully closed.
	E. Loss of NO delivery.	a. If the INOblender® is available, manually ventilate the patient (see INOblender Operation Manual). or b. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual).
3. High NO ₂ Alarm  High NO₂	A. Incomplete system purge.	a. Perform a system purge. See Section 3/ Pre-Use Checkout. 
	B. Ventilator flow stopped.	a. Allow the ventilator gas to flush NO and NO ₂ from the breathing circuit before connecting to the patient.
	C. Two cylinder valves are open.	a. Close one of the cylinder valves.
	D. The NO ₂ alarm limit may be set too low.	a. Make sure the NO ₂ alarm limit is appropriate for the Set NO level.
	E. The NO ₂ calibration may have drifted.	a. Perform a low and high range calibration of the NO ₂ sensor. b. Check calibration sample tee for leaks.
	F. Out of date or the wrong calibration gas was used.	a. Verify the calibration gas expiration date. b. If needed replace the calibration gas and perform a low and high range calibration of the NO ₂ sensor.
	G. The patient circuit setup may be incorrect.	a. Make sure the patient circuit hoses and lengths are correct (see Section 4/ Patient Application of Operation and Maintenance Manual). b. Verify the humidifier chamber is less than 480 mL.
	H. Sample line occlusion. In this case, this alarm may occur with a Sample Line Block alarm.	a. Confirm whether the High NO ₂ alarm occurs concurrently with a sample line block alarm. b. If so, this alarm will clear within 10 seconds after the sample line fault is remedied.
	I. The INOmax® DS _{IR} may have failed.	a. Contact Customer Service. b. Replace the delivery system if in use. c. Do not use the delivery system until serviced.

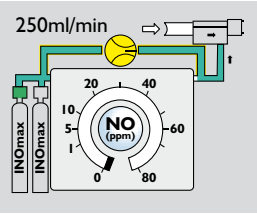
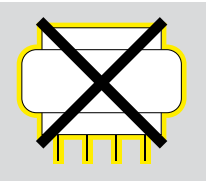
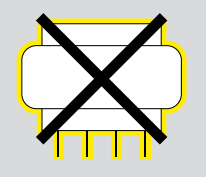
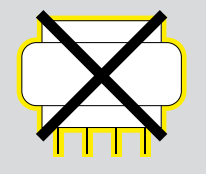
High Priority Alarms

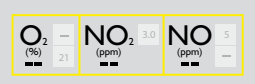
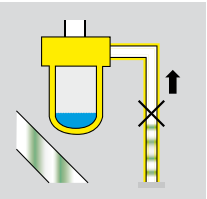
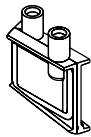
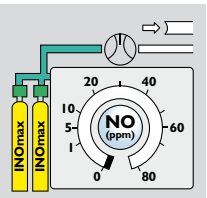
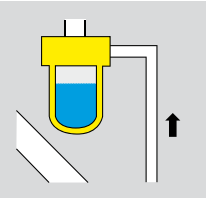
Symptom/Alarm	Possible Cause	Recommended Action
4. High O ₂ Alarm  High O₂	A. The O ₂ alarm setting may be inappropriate. B. The O ₂ calibration may have drifted.	a. Make sure the High O ₂ alarm is set appropriately for the O ₂ setting being used on the ventilator. a. Perform a low and high range calibration of the O ₂ sensor. b. Change the O ₂ sensor if the monitor fails to calibrate. c. Contact Customer Service. d. Replace the delivery system if in use. e. Do not use the delivery system until service.
5. Low O ₂ Alarm  Low O₂	A. The O ₂ concentration setting at the ventilator was reduced. B. The O ₂ alarm setting may be inappropriate. C. The O ₂ sensor may not be properly seated. D. The O ₂ calibration may have drifted.	a. Make sure the O ₂ alarm setting is correct for the setting at the ventilator. a. The INOmax DS _{IR} can dilute the O ₂ concentration set at the ventilator by up to 10%. b. Verify that the alarm is set appropriately for the O ₂ setting being used on the ventilator. a. Make sure the sensors are correctly seated and the sensor cover is fully closed. a. Perform a low and high range calibration of the O ₂ sensor. b. Contact Customer Service.
6. Cylinder Not Detected  or Cylinder Valve Closed 	A. Interference with the Infrared communication link between the INOMAX cylinder and the INOmax DS _{IR} . (Delivery Stopped will occur one hour from the point when communication is lost). B. INOMAX cylinder valve is closed. (Delivery Stopped will occur one hour from the point when the cylinder valve is closed). C. Transport Cap not connected to the INOmeter. (Delivery Stopped will occur one hour from the point when communication is lost). D. INOMAX cylinder not present on the INOmax DS _{IR} . (Delivery Stopped will occur one hour from the point when communication is lost). E. INOmeter may have failed. F. INOmax DS _{IR} infrared cart cable is not connected or has failed.	a. Remove obstruction between the INOMAX cylinder and the INOmax DS _{IR} . b. Move the interfering light or the cart to reduce the high intensity light in the area of the INOmeter®. a. Open INOMAX cylinder valve. a. Attach the Transport Cap to the INOmeter on the INOMAX cylinder. b. Connect the Transport Regulator/Cap Assembly cable to the infrared connector on the back of the INOmax DS _{IR} . a. Load an INOMAX cylinder onto the INOmax DS _{IR} cart. a. Replace the INOMAX cylinder on the INOmax DS _{IR} cart. a. Connect infrared cart cable to the infrared connector on the back of the INOmax DS _{IR} . b. Replace the INOmax DS _{IR} .

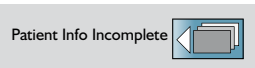
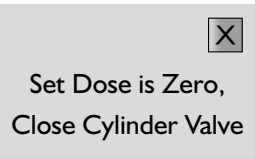


High Priority Alarms		
Symptom/Alarm	Possible Cause	Recommended Action
7. Delivery Failure 	A. Monitored NO levels ≥ 100 ppm or B. The INOmax DS _{IR} has failed.	a. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or b. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual). c. Power the INOmax DS _{IR} to STANDBY and then back ON to reset the delivery system. If this does not work contact Customer Service. d. Replace the delivery system if in use. e. Do not use the delivery system until serviced.
8. Delivery Stopped 	A. Infrared communication link between the INOMAX cylinder and the INOmax DS _{IR} has been lost for one hour. B. INOMAX cylinder is expired and cylinder valve has been open for two minutes. C. INOMAX cylinder is the wrong concentration and cylinder valve has been open for two minutes. D. INOmeter may have failed. E. INOMAX cylinder valve is closed. F. INOMAX cylinder not present on the INOmax DS _{IR} .	a. Remove obstruction between the INOMAX cylinder and the INOmax DS _{IR} . b. Move the interfering light or the cart to reduce the high intensity light in the area of the INOmeter. c. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or d. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual). a. Remove the INOMAX cylinder from the INOmax DS _{IR} cart. b. Connect an INOMAX cylinder to the INOmax DS _{IR} with a valid expiration date. a. Remove the INOMAX cylinder with the wrong concentration from the INOmax DS _{IR} cart. b. Connect an INOMAX cylinder to the INOmax DS _{IR} with a valid concentration. a. Replace the INOMAX cylinder on the INOmax DS _{IR} cart. a. Open INOMAX cylinder valve. b. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or c. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual). a. Load an INOMAX cylinder onto the INOmax DS _{IR} cart. b. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or c. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual).

High Priority Alarms		
Symptom/Alarm	Possible Cause	Recommended Action
9. Drug Past Expiry Date or Drug Concentration Mismatch 	A. INOMAX cylinder is expired (Delivery Stopped will occur two minutes from the point when the cylinder valve is opened).	a. Close the cylinder valve. b. Remove expired INOMAX cylinder from the INOMax DS _{IR} cart. c. Replace the expired INOMAX cylinder on the INOMax DS _{IR} cart.
	B. INOMAX cylinder is the wrong concentration. (Delivery Stopped will occur two minutes from the point when the cylinder valve is opened).	a. Close the cylinder valve. b. Remove the INOMAX cylinder with the wrong concentration from the INOMax DS _{IR} cart. c. Replace the expired INOMAX cylinder with the wrong concentration on the INOMax DS _{IR} cart.
10. Injector Module Fail 	A. The Injector Module electrical cable may be disconnected.	a. Reconnect the Injector Module electrical cable. b. Turn OFF the INOMax DS _{IR} set dose to silence the alarm. c. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or d. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual).
	B. The Injector Module may have failed.	a. Turn OFF the INOMax DS _{IR} set dose to silence the alarm. b. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or c. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual) d. Replace the Injector Module. e. Set the delivered dose and turn OFF the INOblender or the backup mode.
	C. The Injector Module electrical cable may have failed.	a. Turn OFF the INOMax DS _{IR} set dose to silence the alarm. b. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or c. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual). d. Replace the Injector Module electrical cable. e. Reset the delivered dose and turn OFF the INOblender or the backup mode.

High Priority Alarms		
Symptom/Alarm	Possible Cause	Recommended Action
11. Low Battery Alarm 	A. Battery is running low (approximately 10 minutes or less until battery depletion).	a. Check main power indicator.  b. Connect to AC main power source. c. Make sure the power cord is fully inserted into the Power Cord Inlet and that the power cord clamp is secure. d. Check and replace fuse if necessary. e. Contact Customer Service.
12. Low NO/N ₂ Pressure 	A. The NO cylinder supply may be low.	a. Make sure the NO cylinder is turned ON. b. If the high pressure cylinder gauge reads less than 200 psig, change the cylinder. c. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or d. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual).
	B. The supply line may not be connected.	a. If the cylinder gauge reads greater than 200 psig. b. Verify the low pressure hoses are connected correctly to the back of the INOmax DS _{IR} . c. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or d. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual).
13 . Service Required 	A. The INOmax DS _{IR} has failed.	a. If the INOblender is available, manually ventilate the patient (see INOblender Operation Manual). or b. Turn the backup mode ON (see page 29 of Operation and Maintenance Manual). c. Turn the INOmax DS _{IR} to STANDBY and then back ON to rest the delivery system. Please contact Customer Service and replace the delivery system as soon as possible.
14. Set NO and Backup On 	A. The backup mode has been turned ON and the set dose is still set.	a. Turn the INOmax DS _{IR} set dose to zero. b. Correct the reason for initiating the backup mode. c. Turn ON the INOmax DS _{IR} set dose. d. Turn the backup mode OFF. e. If this does not work contact Customer Service.

Low Priority Alarms		
Symptom/Alarm	Possible Cause	Recommended Action
15. Backup On 	A. The backup mode has been turned ON and the set dose is zero.	a. Correct the reason for initiating the backup mode. b. Turn ON the INOmax DS _{IR} set dose. c. Turn the backup mode OFF.
16. Failed NO Sensor 	A. The wrong calibration gas may have been used.	a. Make sure the correct calibration gas is used and ensure sample tubing connections are secure and do not leak.
	B. There is a leak around the sensors.	a. Make sure the sensors are correctly seated with the O-rings and the sensor cover is fully closed.
	C. NO sensor absent or failed.	a. Complete a low calibration first, and then repeat the high calibration. b. Sensor needs to be replaced. c. Replace the delivery system if in use. d. Contact Customer Service.
17. Failed NO ₂ Sensor 	A. The wrong calibration gas may have been used.	a. Make sure the correct calibration gas is used and ensure sample tubing connections are secure and do not leak.
	B. There is a leak around the sensors.	a. Make sure the sensors are correctly seated with the O-rings and the sensor cover is fully closed.
	C. NO ₂ sensor absent or failed.	a. Complete a low calibration first, and then repeat the high calibration. b. Sensor needs to be replaced. c. Replace the delivery system if in use. d. Contact Customer Service.
18. Failed NO ₂ Sensor 	A. The wrong calibration gas may have been used.	a. Make sure the correct calibration gas is used and ensure sample tubing connections are secure and do not leak.
	B. There is a leak around the sensors.	a. Make sure the sensors are correctly seated and the sensor cover is fully closed.
	C. O ₂ sensor absent or failed.	a. Complete a low calibration first, and then repeat the high calibration. b. Sensor needs to be replaced. c. Replace the delivery system if in use. d. Contact Customer Service.

Low Priority Alarms		
Symptom/Alarm	Possible Cause	Recommended Action
19. Monitoring Failure 	A. Monitor is failing to communicate correctly or is reporting a fault.	a. Does not stop delivery of INOMAX to the patient. b. Contact Customer Service.
20. Sample Line/Filter Block 	A. The sample line may be blocked.	a. Make sure the sample inlet line and outlet ports are not obstructed. b. Change the sample line.
	B. The water separator cartridge may be blocked.	a. Replace the water separator cartridge. 
21. Two Cylinders Open 	A. Two cylinder valves are open.	a. Close one of the cylinder valves.
22. Water Trap Bottle Full 	A. The water trap bottle on the side of the INOMAX DS _{IR} is full.	a. Empty the water trap bottle.
	B. Water trap bottle is empty but the message remains in the alarm message box.	a. Remove the water trap bottle and clean the optical sensor level indicator with an alcohol swab.
	C. The INOMAX DS _{IR} may have failed.	a. Contact Customer Service. b. Replace the delivery system if in use.

Indicators		
Symptom/Indicator	Possible Cause	Recommended Action
23. Patient Info Incomplete 	A. Patient identifier has not been entered.	a. Enter patient identifier.
24. Set Dose is Zero, Close Cylinder Valve 	A. The set dose has been set to zero and the INOMAX cylinder valve is still open.	a. Close the INOMAX cylinder valve if treatment has been stopped.
25. Running on Battery 	A. Device is operating on the battery.	a. Connect to AC main power source when available. b. Check main power indicator.  c. Make sure the power cord is fully inserted into the Power Cord Inlet and that the power cord clamp is secure.